Influenza Virus Vaccine Live, Intranasal (LAIV) Spray

Manufacturer MedImmune

Brand Name FluMist®

Age Healthy, non pregnant person 2 years - 49 years of age

Formulation 10 pre-filled single-use sprayers (0.2 ml)

Dosage 0.1 ml in each nostril (thimerosal mercury content = 0 mcg)

Storage Stored in a refrigerator between 2-8°C (35-46°F) **Do Not Freeze**

Injection Site N/A

Route Intranasal only (Do not inject or use a needle)

Needle Size N/A

Administration Should not be administered until 48 hours after cessation of influenza A and/or B antiviral therapy.

Influenza antiviral medications should not be administered for 2 weeks after receipt of FluMist®.

Children age 3yrs-8 yrs may need more than one dose.

Refer to the Seasonal Influenza dosing chart: http://www.kdheks.gov/flu/index.html

Vaccination efforts should begin as soon influenza vaccine is available and continue through the influenza season

Contraindications to FluMist® administration:

- 1. Persons who have experienced an anaphylactic reaction to eggs, egg proteins, gentamicin, gelatin or arginine, or with a life-threatening reaction to a prior dose of influenza vaccine.
- 2. Persons with a history of an egg allergy who have experienced only hives after exposure to eggs should receive TIV
- 3. Persons 2 17 years of age receiving long-term therapy with aspirin or other salicylates, because of the association of Reye syndrome with wild-type influenza infection.

Precautions and Warnings:

- 1. FluMist ® should not be given to any child < 24 months of age because of the increased risk of hospitalizations and wheezing that was observed in clinical trials.
- 2. FluMist ®should not be administered to any individuals with asthma or children < 5 years of age with recurrent wheezing because of the potential for increased risk of wheezing post vaccination unless the potential benefit outweighs the potential risk. If a HCW has told the parent or guardian in the past 12 months their child has asthma or had a wheezing episode in the child's medical record they should not receive FluMist®.
- Persons with chronic medical conditions, including asthma, a recent wheezing episode, reactive airways disease or other chronic pulmonary or cardiovascular conditions, metabolic disease such as diabetes, renal disease, or hemoglobinopathy, such as sickle cell disease.
- 4. History of Guillian-Barre` syndrome within 6 weeks of any prior influenza vaccination. The decision to give Flu Mist should be based on careful consideration of the potential benefits and potential risks.
- 5. Data supporting the safety and effectiveness in immunocompromised individual pregnant women, geriatric adults, or children < 2 yrs is limited.
- 6. Persons who are moderately or severely ill.

Report all adverse reactions to influenza vaccine to the federal Vaccine Adverse Event Reporting System (VAERS) at www.vaers.hhs.gov or (800) 822-7967

Medical Director's Signature:	Effective Date:
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http://www.cdc.gov/mmwr/preview/mmwrhtml/mm6132a3.htm?s_cid=mm6132a3_w	
Drug Insert: http://www.fda.gov/downloads/RiologicsRloadVaccines/Vaccines/ApprovedProducts/UCM123743.pdf	

CDC Influenza website http://www.cdc.gov/flu

KDHE Influenza website http://www.kdheks.gov/flu/index.html